PATENT COOPERATION TREATY

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(PCT Article 36 and Rule 70)

Applicant's or agent's file reference 4 -32610A/HO 61		FOR FURTHER ACTION	See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)			
International application No. PCT/EP 03/08824		International filing date (day/mod 08.08.2003	nth/year)	Priority date (day/month/year) 09.08.2002		
International Patent Classification (IPC) or both national classification and IPC CO7D277/68						
Applicant NOVARTIS AG et al.						
1. This inte	. This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.					
2. This RE	2. This REPORT consists of a total of 5 sheets, including this cover sheet.					
be	This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).					
These a	These annexes consist of a total of sheets.					
3. This rep	ort contains indications re	lating to the following items:				
ı ⊠	Basis of the opinion					
11 🗆	•					
🗆	Non-establishment of	opinion with regard to novelty,	inventive step a	and industrial applicability		
	•					
V ⊠		under Rule 66.2(a)(ii) with rega ions supporting such statemer		ventive step or industrial applicability;		
VI 🗆	Certain documents cit	ed				
VII 🗆	Certain defects in the	international application				
VIII □	Certain observations of	on the international application	•			
Date of submission of the demand			of completion of th	als report		
16.02.2004			1.2004			
Name and mailing address of the international preliminary examining authority:			rized Officer	Alienes Palanteny.		
European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 epmu d Fax: +49 89 2399 - 4465			lon, L hone No. +49 89 :	2399-7078		

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No.

PCT/EP 03/08824

I. Basis of the report

1. With regard to the **elements** of the international application (Replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report since they do not contain amendments (Rules 70.16 and 70.17)):

	Description, Pages						
	1-6		as originally filed				
		-	as engineary med				
	Cla	laims, Numbers					
	1-1	4	as originally filed				
2.	With lang	h regard to the langu guage in which the int	age, all the elements marked above were available or furnished to this Authority in the ternational application was filed, unless otherwise indicated under this item.				
	The	ese elements were av	ailable or furnished to this Authority in the following language: , which is:				
		the language of a translation furnished for the purposes of the international search (under Rule 23.1(b)).					
		the language of publ	lication of the international application (under Rule 48.3(b)).				
		the language of a tra Rule 55.2 and/or 55.	anslation furnished for the purposes of international preliminary examination (under 3).				
3.	With regard to any nucleotide and/or amino acid sequence disclosed in the international application, the international preliminary examination was carried out on the basis of the sequence listing:						
		contained in the inte	rnational application in written form.				
		filed together with the international application in computer readable form.					
		furnished subsequently to this Authority in written form.					
		furnished subsequently to this Authority in computer readable form.					
		The statement that to in the international a	statement that the subsequently furnished written sequence listing does not go beyond the disclosic international application as filed has been furnished.				
		The statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished.					
4.	The	amendments have re	esulted in the cancellation of:				
		the description,	pages:				
		the claims,	Nos.:				
		the drawings,	sheets:				
5.	×	This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)).					
		(Any replacement sheet containing such amendments must be referred to under item 1 and annexed to this report.)					
	see separate sheet						

Form PCT/IPEA/409 (January 2004)

6. Additional observations, if necessary:





International application No.

PCT/EP 03/08824

V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)

Yes: Claims

7-8,14

No: Claims

1-6, 9-13

Inventive step (IS)

Yes: Claims

No: Claims

7-8,14

Industrial applicability (IA)

Yes: Claims

1-14

No: Claims

2. Citations and explanations

see separate sheet

Re Item I

Basis of the report

This opinion has been established as if the amendments dated 05.10.2004 had not been made since they have been considered to go beyond the disclosure as filed (rule 70.2(c)) for the following reasons:

- a disclaimer can only restore novelty against an accidental anticipation
- a disclaimer should not remove more subject-matter than is necessary (i.e. compound S1319 disclosed in D1 and D2).

Re Item V

Reasoned statement with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

Reference is made to the following documents:

D1: INFLAMMATION RESEARCH (2000),49(2), 86-94

D2: BIOORGANIC & MEDICINAL CHEMISTRY LETTERS, 1999, vol. 9, no. 10, pages

1361-1364

D3: US-A-5 648 370

1). Claims 1 to 6 and 9 to 13 lack novelty with respect to the compounds named S1319 disclosed in D1 and D2 (Article 33(2) PCT).

The subject-matter of claims 7 to 8 is novel over D1-D3 on account of the moiety R¹ArR² of the claimed derivatives of formula II.

2). The technical problem underlying the present application is seen in the provision of derivatives having beta 2 adrenoreceptor agonistic activity.

D3 which discloses benzothiazolone derivatives having that activity is considered to represent the closest prior art (see D3, formula I, column 5, lines 45, column 6, lines 22 to 24).

The claimed derivatives differ from those of D3 only on account of the presence of the hydroxy group on the ethylene linking group between the amino and the benzothiazolone moiety.

From the cited documents D1 and D2 it appears that the benzothiazolone derivatives

can further be substituted with a hydroxy substituent in that position with the retention of the activity of the resulting products.

Accordingly, the skilled man faced with the above mentioned technical problem and being aware that this structural modification would not impair the beta 2 adrenoreceptor agonistic activity of the derivatives would have considered the claimed derivatives of claim 7 as obvious solutions.

Inventive step could however be acknowledged if the present compounds were shown to exhibit unexpected properties over D1-D3 (e.g. a rapid onset of action, long duration of action etc.). Said properties establishing an inventive step should be clearly established for example with comparative tests and should extend to the whole claimed scope.